Section C 510(k) Summary (21 CFR 807.92)

510(K) Summary

"This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirements of SMDA 1990 and 21 CFR 807.92."

"The assigned 510(k) number is: K130101 "

Premarket Notification [510(k)] Summary

NOV 08 2013

Submitter's name :	Bytech Dongtai Co.,Ltd.
Submitter's address :	Xing Yuan Industrial Park, Tang Yang Town, Dong Tai City, JiangSu, 224200 China
Phone number :	0086-515-85655500
Fax number :	0086-515-85655500
Name of contact person:	Mr.Wang Cheng
Date the summary was prepared:	2013-09-29
Device Name:	Powder Free Vinyl Patient Examination Gloves, Clear (non-colored)
Proprietary/Trade name:	**"Bytech"**
Common Name:	Exam gloves
Classification Name:	Patient examination glove
Device Classification:	I
Regulation Number:	21 CFR 880.6250
Panel:	General Hospital (80)
Product Code:	LYZ

Class I* Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) that meets all of the requirements of ASTM D 5250-06 (Reapproved 2011).

Predicate device: Powder-Free Vinyl Patient Examination Glove (Non-colored) Zhang Jia Gang Fengyuan Plastic Product Co., Ltd. K091663.

Device Description: Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) are disposable devices which made of PVC material ,intended for medical purpose that worn on examiner's hand or finger to prevent contamination between patient and examiner and they meets all of the requirements of ASTM standard D 5250-06 (Reapproved 2011).

Device Intended Use (Indication for use): Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

A summary of the technological characteristics of new device compared to the predicate device.

The Powder Free Vinyl Patient Examination Gloves, Clear (non-colored), non sterile are summarized with the following technological characteristics compared to ASTM or equivalent standard.

Characteristics	Standard	Device performance
Dimension	ASTM standard D 5250-06(Reapproved 2011).	Meets
Physical Properties	ASTM standard D 5250-06(Reapproved 2011).	Meets
Freedom from pinholes	21 CFR 800.20	Meets
Powder Residual	ASTM standard D 5250-06 (Reapproved 2011). and D6124-06(Reapproved 2011).	Meets <2mg/glove
Biocompatability	Primary Skin Irritation in rabbits ISO 10993-10	Passes The test article was non irritant.
	Dermal sensitization in the guinea pig ISO 10993-10	Passes The test article was non sensitizer.

A brief discussion of the nonclinical submitted, reference, or relied on in the premarket notification submission for a determination of substantial equivalence .

Powder Free Vinyl Patient Examination Gloves, Clear (non-colored), meet requirements per ASTM D5250-06 (Reapproved 2011), per ASTM D6124-06(Reapproved 2011), per 21 CFR 800.20 and ISO 10993-10: 2002/Amd.1:2006.
The performance test data of the nonclinical tests that support a determination of substantial equivalent is the same as mentioned immediately above.

Features & Description	Predicate Device	Medical Glove Guidance Manual(1661)	Subject Device	Result of Comparison
Company	Zhang Jia Gang Fengyuan Plastic Product Co.Ltd.		Jiangsu Toptouch Glove Co., Ltd.	--
510(K) Number	K091663		--	
Product name	Powder Free Vinyl Patient Examination Gloves, Clear (Non-colored)		Powdered Free Vinyl Patient Examination Gloves, Clear (non-colored)	same
Product Code	LYZ	LYZ	LYZ	same
Size	Small/ Medium/ Large/X large		Small/ Medium/ Large/X large	same
Intend for use	Powder free Vinyl Patient Examination Gloves, Clear(Non-colored)is a disposable device	Powder Free Examination Gloves: A powder-free patient examination glove is a disposable device	Powder free Vinyl Patient Examination Gloves, Clear (Non-colored) is a disposable device	Substantially equivalent

	intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	
Device Description and Specifications	Meets ASTM D5250-06 (Reapproved 2011)	If vinyl: Do the vinyl examination gloves meet all the current specifications listed under ASTM Specification D5250 or an equivalent consensus standard?	Meets ASTM D5250-06 (Reapproved 2011)	Substantially equivalent
Dimensions --Length	Meets ASTM D5250 -06 (Reapproved 2011) ≥230mm min	ASTM D5250	230mm min for all sizes	Substantially equivalent
Dimensions -- Width	Meets ASTM D5250-06 (Reapproved 2011) Small 80-90 mm Medium 90-100mm Large 100-110mm Xlarge 110-120 mm	ASTM D5250	 Small 80-85 mm Medium 95-100mm Large 102-108mm X large 113-118 mm	Substantially equivalent
Dimensions --Thickness	Meets ASTM D5250-06 (Reapproved 2011) Finger 0.05mm min. Palm 0.08mm min.		Thickness (mm) min. Finger 0.09-0.12 Palm 0.09-0.12	Substantially equivalent
Physical Properties	Meets ASTM D 5250-06 (Reapproved 2011) Before aging/after aging Elongation ≥300% Tensile Strength≥ 14MPa	ASTM D5250	 Before aging/after aging Elongation :390-420% Tensile Strength:15-20 MPa	Substantially equivalent
Freedom from Pinholes	Meets • 21 CFR 800.20 • ASTM D5250-06 (Reapproved 2011) • ASTM D 5151-06 (Reapproved 2011)	21 CFR 800.20 ASTM D5250 ASTM D 5151	Meets ASTM D5151-06 (Reapproved 2011) Holes at Inspection Level I AQL2.5	Substantially equivalent
Residual Powder	Meets ASTM D 6124-06 (Reapproved 2011) below 2mg of residual powder	ASTM D 6124	Meets ASTM D 6124-06 (Reapproved 2011) Results generated values below 2mg of residual powder	Substantially equivalent
Materials used to fabricate the devices	PVC	If the glove is made of a polymer or other type of material. identify the material.	PVC	Substantially equivalent
Dusting or Donning	PU	If a donning lubricant is used, state the	PU	Substantially equivalent

Powder:		composition and include biocompatibility data for the lubricant in an identified attachment; also state the name, manufacturer, and address below		
Dusting or Donning Powder: name	PU	Lubricant Generic Name/ Lubricant Brand Name	Surface Coating Agent	Substantially equivalent
Compare performance data supporting substantial equivalence	Meets ASTM D5151-06 (Reapproved 2011) ASTM D5250-06 (Reapproved 2011) ASTM D6124-06 (Reapproved 2011)	At this time FDA recognizes the following standards: Patient Examination Gloves(PVC)ASTM D5151(Detection of Holes in Medical Gloves)ASTM D6124(Residual Powder on Medical Gloves)ASTM D5250(Poly(vinyl chloride) Gloves)	Meets ASTM D5151-06 (Reapproved 2011) ASTM D5250-06 (Reapproved 2011) ASTM D6124-06 (Reapproved 2011)	Substantially equivalent
Single Patient Use	Single Patient Use	Single Patient Use	Single Patient Use	Substantially equivalent
Biocompatibility	SKIN IRRITATION DERMAL and SENSITIZATION STUDIES Meets ISO 10993-10:2002/Amd.1: 2006	SKIN IRRITATION DERMAL and SENSITIZATION STUDIES ISO 10993-10	The test article was non irritant and non sensitizer. SKIN IRRITATION DERMAL and SENSITIZATION STUDIES Meets ISO 10993-10:2002 /Amd.1:2006	Substantially equivalent
Labeling for the legally marketed device to which substantial equivalence is claimed.	-Powder Free -Patient Examination Glove -Single Use Only - Manufactured For: - Lot	Chapter 4	-Powder Free -Patient Examination Glove -Single Use Only - Manufactured For: - Lot	Substantially equivalent

A brief discussion of the clinical submitted, reference, or relied on in the premarket notification submission for a determination of substantial equivalence .

Clinical data is not needed for gloves or for most devices cleared by the 510(k) process.

The conclusions

It can be concluded that the Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) meet the ASTM standard or equivalent standard and FDA requirements for waterleak test on pinhole AQL., meet labeling claims.

It can be concluded that the Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) is as safe, as effective, and performs as well as the predicate device, Powder-Free Vinyl Patient Examination Glove (Non-colored) Zhang Jia Gang Fengyuan Plastic Product Co., Ltd. K091663.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
10903 New Hampshire Avenue
Document Control Center – WO66-G609
Silver Spring, MD 20993-0002

November 8, 2013

Bytech Dongtai Company, Limited
Mr. Wang Cheng
Quality Department Manager
Xing Yuan Industrial Park, Tang Yang Town
Dong Tai City, JiangSue
CHINA 224200

Re: K130101
Trade/Device Name: Powder Free Vinyl Patient Examination Gloves, Clear (non-colored)
Regulation Number: 21 CFR 880.6250
Regulation Name: Patient Examination Glove
Regulatory Class: I
Product Code: LYZ
Dated: September 29, 2013
Received: October 2, 2013

Dear Mr. Cheng:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

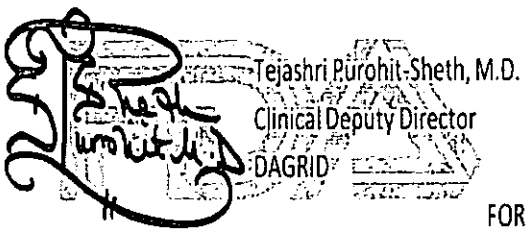
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm.

Sincerely yours,



FOR

Kwame Ulmer M.S.
Acting Division Director
Division of Anesthesiology, General Hospital,
Respiratory, Infection Control and
Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120
Expiration Date: December 31, 2013
See PRA Statement on last page.

510(k) Number *(if known)*
K130101

Device Name
Powder Free Vinyl Patient Examination Gloves, Clear (non-colored)

Indications for Use *(Describe)*
Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) is a non-sterile disposable device intended for medical purpose that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

Type of Use *(Select one or both, as applicable)*

☐ Prescription Use (Part 21 CFR 801 Subpart D) ☒ Over-The-Counter Use (21 CFR 801 Subpart C)

PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON A SEPARATE PAGE IF NEEDED.

FOR FDA USE ONLY

Concurrence of Center for Devices and Radiological Health (CDRH) *(Signature)*

Elizabeth F. Claverie -S
2013.11.07 21:50:02 -05'00'